Attorney Docket No. 010180.00051

The listing of claims will replace all prior versions, and listings, of claims in the application: Listing of Claims:

1. (Currently Amended) The use of a A compound of formula (I), or a salt, N-oxide, hydrate, or solvate thereof, in the preparation of a composition for inhibition of HSP90 activity in vitro or in vivo:

$$R_2$$
 R_3 R_4 (I)

wherein

R₂ is a group of formula (IA):

$$-(Ar^{1})_{m}-(Alk^{1})_{p}-(Z)_{r}-(Alk^{2})_{s}-Q$$
 (IA)

wherein in any compatible combination

Ar1 is an optionally substituted aryl or heteroaryl radical,

Alk¹ and Alk² are optionally substituted divalent C_1 - C_3 alkylene or C_2 - C_3 alkenylene radicals,

m, p, r and s are independently 0 or 1,

wherein R^A is hydrogen or C_1 - C_6 alkyl, and

Q is hydrogen or an optionally substituted carbocyclic or heterocyclic radical;

 R_3 is hydrogen, an optional substituent, or an optionally substituted (C_1C_6)alkyl, aryl or heteroaryl radical; and

R₄ is a carboxylic ester, carboxamide or sulfonamide group.

- 2. (Currently Amended) The use compound as claimed in claim 1 wherein m is 1, each of p, r and s is 0, and Q is hydrogen.
- 3. (Currently Amended) The use-compound as claimed in claim 2 wherein R₂ is optionally substituted phenyl, 2- or 3-thienyl, 2- or 3-furanyl, or 2-, 3- or 4-pyridinyl.
- 4. (Currently Amended) The use compound as claimed in claim 2 wherein R₂ is phenyl, optionally substituted by methyl, ethyl, n- or isopropyl, methoxy, ethoxy, isopropoxy, chloro, or bromo.
- 5. (Currently Amended) The use-compound as claimed in claim 3 wherein the optional substituent is in the 4-position of the phenyl ring.
- 6. (Currently Amended) The <u>use-compound</u> as claimed in claim 1 wherein m is 1, and p, r and s are 0, and Q is an optionally substituted carbocyclic or heterocyclic ring.
- 7. (Currently Amended) The use compound as claimed in claim 1 wherein Ar¹ is a phenyl or pyridyl ring.
- 8. (Currently Amended) The use-compound as claimed in any of the preceding claims claim 1 wherein R₃ is amino (NH₂).
- 9. (Currently Amended) The use-compound as claimed in any of the preceding claims claim 1 wherein R₄ is a carboxamide group of formula -CONR^B(Alk)_nR^A wherein

Alk is a divalent alkylene, alkenylene or alkynylene radical, for example a -CH₂-, -CH₂CH₂-, -CH₂CH₂-, -CH₂CH-CH-, or -CH₂CCCH₂-radical, and the Alk radical may be optionally substituted,

n is 0 or 1,

R^B is hydrogen or a C₁-C₆ alkyl or C₂-C₆ alkenyl group, for example methyl, ethyl, nor iso propyl, or allyl,

R^A is hydroxy or optionally substituted carbocyclic, for example hydroxy and/or ehloro-substituted phenyl and 3,4 methylenedioxyphenyl; or heterocyclyl, for example pyridyl, furyl, thienyl, N-piperazinyl, or N-morpholinyl any of which heterocyclic rings may be substituted, or R^A and R^B taken together with the nitrogen to which they are attached form an N-heterocyclic ring which may optionally contain one or more additional hetero atoms selected from O, Sand N, and which may optionally be substituted on one or more ring C or N atoms, examples of such N-heterocyclic rings including morpholino, piperidinyl, piperazinyl and N-phenylpiperazinyl.

- 10. (Currently Amended) The use-compound as claimed in any of claims 1 to 8 claim 1 wherein R_4 is a carboxylic ester group of formula -COOR^C wherein R^C is a C_1 - C_6 alkyl or C_2 - C_6 alkenyl group, or an optionally substituted aryl or heteroaryl group, or an optionally substituted aryl(C_1 - C_6 alkyl)- or heteroaryl(C_1 - C_6 alkyl)- group or an optionally substituted cycloalkyl group.
- 11. (Currently Amended) The use-compound as claimed in any of elaims 1 to 8 claim 1 wherein R_4 is a carboxylic ester group of formula -COOR^C wherein R^C is optionally substituted methyl, ethyl, n- or iso-propyl, allyl, phenyl, pyridyl, thiazolyl, benzyl, pyridylmethyl, cyclopentyl or cyclohexyl.
- 12. (Currently Amended) A method of treatment of diseases or conditions mediated by excessive or inappropriate HSP90 activity in mammals which method comprises administering to the mammal an amount of a compound as defined in any of claims 1 to 11 claim 1 effective to inhibit said HSP90 activity.
- 13. (Currently Amended) The use as claimed in claim 11 or a method as claimed claim 12 for the treatment of cancer.

- 14. (Currently Amended) The use as claimed in claim-11 or a method as claimed claim 12 for immunosuppression or the treatment of inflammatory diseases such as rheumatoid arthritis, asthma, multiple selerosis, Type I diabetes, lupus, psoriasis and inflammatory bowel disease; or cystic fibrosis angiogenesis-related disease such as diabetic retinopathy, haemangiomas, and endometriosis; or for protection of normal cells against chemotherapy-induced toxicity; or diseases where failure to undergo apoptosis is an underlying factor; or protection from hypoxia-ischemic injury due to elevation of Hsp70 in the heart and brain; scrapie/CJD, Huntingdon's or Alzheimer's disease.
- 15. (Currently Amended) A pharmaceutical or veterinary composition comprising a compound of formula (I) as specified in any of claims 1 to 11, claim 1 together with a pharmaceutically or veterinarily acceptable carrier.
- 16. (New) The compound of claim 9 wherein Alk a -CH₂-, -CH₂CH₂-, -CH₂-, -
- 17. (New) The compound of claim 9 wherein R^B methyl, ethyl, n- or iso-propyl, or allyl.
- 18. (New) The compound of claim 9 wherein R^A is hydroxy, chloro-substituted phenyl, or 3,4 methylenedioxyphenyl; or pyridyl, furyl, thienyl, N-piperazinyl, or N-morpholinyl any of which heterocyclic rings may be substituted.
- 19. (New) The compound of claim 9 wherein R^A and R^B taken together with the nitrogen to which they are attached form morpholino, piperidinyl, piperazinyl or N-phenylpiperazinyl.
- 20. (New) The method as claimed claim 14 for immunosuppression or the treatment of rheumatoid arthritis, asthma, multiple sclerosis, Type I diabetes, lupus, psoriasis, inflammatory bowel disease, diabetic retinopathy, haemangiomas, or endometriosis.